

Amendments to the Claims:

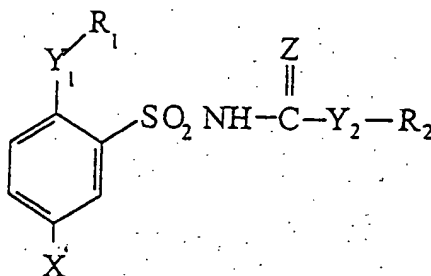
The listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

Claim 1—22 (cancelled)

Claim 23 (new): Benzene-sulphonamide derivatives having the formula (I):

(I)



in which:

X represents a nitro, cyano, or halogen group;

$Y_1$  represents a secondary or tertiary amino group, or a sulphur;

$Y_2$  represents a NH group, or a nitrogen atom in a saturated or unsaturated heterocyclic group having 5 to 7 ring members;

Z represents oxygen, sulphur, -N-CN or -CH-NO<sub>2</sub>; and

$R_1$  and  $R_2$ , which can be identical or different, represent each independently a saturated or unsaturated radio-labeled linear or branched alkyl group with 2 to 12 carbon atoms, a saturated or unsaturated radio-labeled alicyclic group with 3 to 12 carbon atoms, an aryl group substituted or not by one or several alkyl groups in C<sub>1</sub> – C<sub>4</sub>, nitro, cyano, trifluoromethyl, carboxy and halogen groups, or an arylalkyl group

or  $R_1$  and  $Y_1$ , and/or,  $R_2$  and  $Y_2$  form a saturated or unsaturated heterocyclic group having 5 to 7 ring members of which at least one is oxygen or nitrogen

with the exception of compounds for which X is a nitro group,  $Y_1$  represents a secondary amino group (-NH-),  $Y_2$  represents a NH group, Z represents an oxygen,  $R_2$

represents an isopropyl and R<sub>1</sub> represents an element selected from a group consisting of m-toluy, phenyl and cyclootyl, and with the exception of N-[(2-cyclootylamino-5-cyanobenzene)sulfonyl] N'-isopropyl urea.

Claim 24 (new): The derivative according to claim 23, characterized in that X is an element selected from a group consisting of nitro, cyano, bromo and iodine group.

Claim 25 (new): The derivative according to claim 23, characterized in that Y<sub>1</sub> represents a NH group and Y<sub>2</sub> represents a NH group or an oxygen atom.

B<sup>2</sup>  
Claim 26 (new): The derivative according to claim 23, characterized in that R<sub>1</sub> and R<sub>2</sub> represent each independently an ethyl, butyl, tert-butyl, propyl, isopropyl, pentyl, hexyl, heptyl, octyl, decyl, amyl, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl, cyclododecyl, 2-cyclohexenyl, m-toluy, o-toluy, p-toluy, phenyl, allyl, adamantyl, norbornyl, 3-carboxyphenyl, 2,3-dimehtylphenyl, 2,4-dimethylphenyl, 2,5-dimethylphenyl, 2,6-dimethylphenyl, 3,4-dimethylphenyl, 3,5-dimethylphenyl, 2,4,6-trimethylphenyl, furfuryl, benzyl or 1-phenylethyl group.

Claim 27 (new): The derivative according to claim 23, characterized in that R<sub>2</sub> and Y<sub>2</sub> form a homopiperidinyl group.

Claim 28 (new): The derivative according to claim 23, characterized in that R<sub>1</sub> and Y<sub>1</sub> form a morpholinyl or homopiperidinyl group.

Claim 29 (new): The derivative according to claim 23, characterized in that it is constituted by a salt selected from a group consisting of sodium salts, the potassium salts or organic acid salts.

Claim 30 (new): The derivative according to claim 29, characterized in that it is chosen in a group having:

N-[(2-cyclohexylamino-5-nitrobenzene)sulfonyl]N'-tert-butyl urea,

N-cyano-N'-[(2-cyclohexylamino-5-nitrobenzene)sulfonyl]homopiperidinoamidine,  
N-[(2-cyclohexylamino-5-nitrobenzene)sulfonyl]N'-cyclohexyl thiourea, and  
N-[(cyclohexen-2-yl-5-iodobenzene)sulfonyl]N'-pentyl urea.

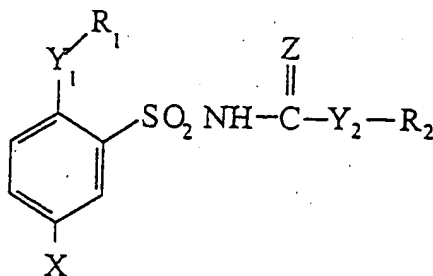
Claim 31 (new): A pharmaceutical composition, characterized in that it includes the benzene sulphonamide derivative according to claim 23 in mixture with an acceptable pharmaceutical excipient.

B<sup>2</sup>  
Claim 32 (new): A method for producing a medication for treatment of an illness involving a thromboxan A<sub>2</sub>, including cardio-vascular and blood, pulmonary, reproduction and renal diseases, which comprises utilizing the derivatives according to claim 23.

Claim 33 (new): A method for binding to a thromboxan A<sub>2</sub> receptor, which comprises utilizing the derivative according to claim 23.

Claim 34 (new): Benzene-sulphonamide derivatives having the formula (I):

(I)



in which:

X represents a nitro, cyano, or radio-labeled halogen group;

Y<sub>1</sub> represents a secondary or tertiary amino group, or a sulphur;

Y<sub>2</sub> represents a NH group, or a nitrogen atom in a saturated or unsaturated heterocyclic group having 5 to 7 ring members;

Z represents oxygen, sulphur, -N-CN or -CH-NO<sub>2</sub>; and

R<sub>1</sub> and R<sub>2</sub>, which can be identical or different, represent each independently a saturated or unsaturated radio-labeled linear or branched alkyl group with 2 to 12 carbon

atoms, a saturated or unsaturated radio-labeled alicyclic group with 3 to 12 carbon atoms, an aryl group substituted or not by one or several alkyl groups in C<sub>1</sub> – C<sub>4</sub>, nitro, cyano, trifluoromethyl, carboxy and halogen groups, or an arylalkyl group

or R<sub>1</sub> and Y<sub>1</sub>, and/or, R<sub>2</sub> and Y<sub>2</sub> form a saturated or unsaturated heterocyclic group having 5 to 7 ring members of which at least one is oxygen or nitrogen

with the exception of compounds for which X is a nitro group, Y<sub>1</sub> represents a secondary amino group (-NH-), Y<sub>2</sub> represents a NH group, Z represents an oxygen, R<sub>2</sub> represents an isopropyl and R<sub>1</sub> represents an element selected from a group consisting of m-toluy, phenyl and cyclootyl, and with the exception of N-[(2-cyclootylamino-5-cyanobenzene)sulfonyl] N'-isopropyl urea.

B<sup>2</sup>

Claim 35 (new): The derivative according to claim 34, characterized in that X is an element selected from a group consisting of nitro, cyano, bromo and iodine group.

Claim 36 (new): The derivative according to claim 34, characterized in that Y<sub>1</sub> represents a NH group and Y<sub>2</sub> represents a NH group or an oxygen atom.

Claim 37 (new): The derivative according to claim 34, characterized in that R<sub>1</sub> and R<sub>2</sub> represent each independently an ethyl, butyl, tert-butyl, propyl, isopropyl, pentyl, hexyl, heptyl, octyl, decyl, amyl, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl, cyclododecyl, 2-cyclohexenyl, m-toluy, o-toluy, p-toluy, phenyl, allyl, adamantyl, norbornyl, 3-carboxyphenyl, 2,3-dimethylphenyl, 2,4-dimethylphenyl, 2,5-dimethylphenyl, 2,6-dimethylphenyl, 3,4-dimethylphenyl, 3,5-dimethylphenyl, 2,4,6-trimethylphenyl, furfuryl, benzyl or 1-phenylethyl group.

Claim 38 (new): The derivative according to claim 34, characterized in that R<sub>2</sub> and Y<sub>2</sub> form a homopiperidinyl group.

Claim 39 (new): The derivative according to claim 34, characterized in that R<sub>1</sub> and Y<sub>1</sub> form a morpholinyl or homopiperidinyl group.

Claim 40 (new): The derivative according to claim 34, characterized in that it is constituted by a salt selected from a group consisting of sodium salts, the potassium salts or organic acid salts.

Claim 41 (new): The derivative according to claim 40, characterized in that it is chosen in a group having:

N-[(2-cyclohexylamino-5-nitrobenzene)sulfonyl]N'-tert-butyl urea,

N-cyano-N'-[(2-cyclohexylamino-5-nitrobenzene)sulfonyl]homopiperidinoamidine,

N-[(2-cyclohexylamino-5-nitrobenzene)sulfonyl]N'-cyclohexyl thiourea, and

N-[(cyclohexen-2-yl-5-iodobenzene)sulfonyl]N'-pentyl urea.

Claim 42 (new): A pharmaceutical composition, characterized in that it includes the benzene sulphonamide derivative according to claim 34 in mixture with an acceptable pharmaceutical excipient.

Claim 43 (new): A method for producing a medication for treatment of an illness involving a thromboxan A<sub>2</sub>, including cardio-vascular and blood, pulmonary, reproduction and renal diseases, which comprises utilizing the derivatives according to claim 34.

Claim 44 (new): A method for binding to a thromboxan A<sub>2</sub> receptor, which comprises utilizing the radio-labeled derivative according to claim 34.

---